

California Drug Recall Information



Recall Name

Nexus Pharmaceuticals Recalls Benztropine Mesylate Injection Due to Visible Particulates in the Vials

Recall Date	Product Description	Recalling Firm	Recall Reason
8/01/13	Benztropine Mesylate Injection, USP 2 mg/2 mL (1 mg/mL), in 2 mL single dose vials NDC #14789-300-02	Distributed by: Nexus Pharmaceuticals, Inc. Vernon Hills, IL	Potential presence of visible particulate matter in the vials.
Recall Class	Product Identification	Distribution	Affected Dates
N/A	Lot Numbers: • 030712, expires 03/2014 • 112911, expires 11/2013 Product Labels	CA, nationwide	Distributed from: 12/15/2011 to 05/17/12

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

http://www.fda.gov/Safety/Recalls/ucm363697.htm

Related recall: http://www.cdph.ca.gov/certlic/manfprocdistrib/Documents/fdbDrFK2n.pdf